

Filovirus Animal Model Workshop

Vaccine Development Panel Discussion

September 11, 2007

Ed Nuzum, NIAID, Chair

Tom Ksiazek, CDC

Alan Schmaljohn, University of Maryland, Med. School

Tom Geisbert, NIAID

Nancy Sullivan, NIAID

Questions

- 1. What is/are the most <u>appropriate animal species</u> to use as models for filoviruses?
- 2. Is there a specific <u>strain</u>, <u>or isolate</u> of virus(es) that should be used in animal studies and what <u>challenge dose</u> should be used if the goal is to protect against a potential bioterrorist release?
- 3. What <u>challenge route/s</u> should be used if the purpose is to develop a vaccine to protect against a bioterrorist attack?
- 4. What types of studies should be required to develop <u>correlates of immunity/protection</u> for filovirus vaccines in humans? Specifically, what <u>immune responses</u> should be examined in animals and what functional <u>assays</u> need to be established and validated?



Filovirus Animal Model Workshop

Therapeutic Panel Discussion

September 12, 2007

Robert Johnson, NIAID, Chair Tony Sanchez, CDC Sina Bavari, USAMRIID Mike Bray, NIAID Doug Reed, USAMRIID

Questions

- 1. What are the similarities or differences in considering developing appropriate animal models for therapeutic countermeasures as compared to that for the vaccines?
- 2. What initial clinical symptoms should be focused on to identify potentially relevant ranges and triggers for timing of treatment initiation for development of therapeutic animal models?
- 3. What types of animal studies may contribute to assessment of antiviral interventions in a post-exposure situation, including approaches to assessment of post-exposure prophylaxis and approaches to assessment of treatment of established illness?
- 4. What are some of the different factors that may be relevant to interventions that focus on inhibition of viral replication as compared to interventions that focus on modification of host responses?
- 5. What approaches can be used to optimize information collection if outbreaks occur at various times during the development sequence for a candidate product?